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RHYTHM DISORDERS AND ELECTROPHYSIOLOGY

CLINICAL CASE SERIES

Safety, Efficacy, and 2-Year Outcomes of Cardiovascular Magnetic Resonance-Guided Ablation of Isthmus-Dependent Atrial Flutter



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ABSTRACT

BACKGROUND Ablation of cavotricuspid isthmus (CTI)-dependent atrial flutter with interventional cardiovascular magnetic resonance (CMR) has demonstrated its efficacy and safety in the initial phase for achieving bidirectional block. However, limited long-term follow-up data exist for these patients.

CASE SUMMARY Thirteen consecutive patients were included between 2021 and 2022. CMR-guided catheter ablation of CTI-dependent atrial flutter was completed successfully in 11 patients (85%). CTI ablation had to be completed in a conventional electrophysiology laboratory for 2 patients: one patient required general anesthesia due to pain during CTI ablation, and the other deteriorated into atrial fibrillation during ablation shots. Among 11 patients with procedural success, 2 (18%) were lost to follow-up after the ablation and 9 (82%) were free of recurrent isthmus-dependent atrial flutter at a median follow-up of 27 months (IQR: 24-29).

CONCLUSIONS Interventional CMR-guided ablation of CTI-dependent atrial flutter demonstrates a good safety profile and efficacy at 2-year follow-up. (JACC Case Rep. 2025;30:104184) © 2025 The Authors. Published by Elsevier on behalf of the American College of Cardiology Foundation. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

BACKGROUND AND AIMS

Catheter ablation in an electrophysiology laboratory with the guidance of fluoroscopy is the first-line therapy for patients presenting with cavotricuspid isthmus (CTI)-dependent atrial flutter. Recently, several studies reported the feasibility of CTI-

TAKE-HOME MESSAGES

- iCMR-guided flutter ablation is feasible, safe and accurate under tailored conscious sedation and in-room monitoring.
- The study shows the accuracy of iCMR-guided flutter ablation over 2-year follow-up.

From the Institut Cardiovasculaire Paris Sud, Hôpital Privé Jacques Cartier, Ramsay Santé, Massy, France. The authors attest they are in compliance with human studies committees and animal welfare regulations of the authors' institutions and Food and Drug Administration guidelines, including patient consent where appropriate. For more information, visit the Author Center.

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ABBREVIATIONS AND ACRONYMS

CMR = cardiovascular magnetic resonance

CTI = cavotricuspid isthmus

ECG = electrocardiogram

iCMR = interventional cardiovascular magnetic resonance

RF = radiofrequency

VT = ventricular tachycardia

dependent atrial flutter ablation under interventional cardiovascular magnetic resonance (iCMR) guidance. Indeed, the development of specific catheters has allowed active catheter tracking in cardiac chambers in real time under iCMR. Compared with fluoroscopy, the advantage of iCMR is the real-time visualization of the tip of the catheter along with the anatomy of the inferior vena cava and cardiac chambers, thereby overcoming some anatomic variations such as a ridge at the junction between the inferior

vena cava and the right atrium. Bijovet et al³ recently showed in a small sample of 15 patients that T1 mapping by cardiovascular magnetic resonance (CMR) could identify regions of acute reconnection directly after ablation. Some studies have demonstrated the efficacy and safety of the procedure in the initial phase for achieving bidirectional block;¹⁻³ however, limited long-term follow-up data exist for these patients.

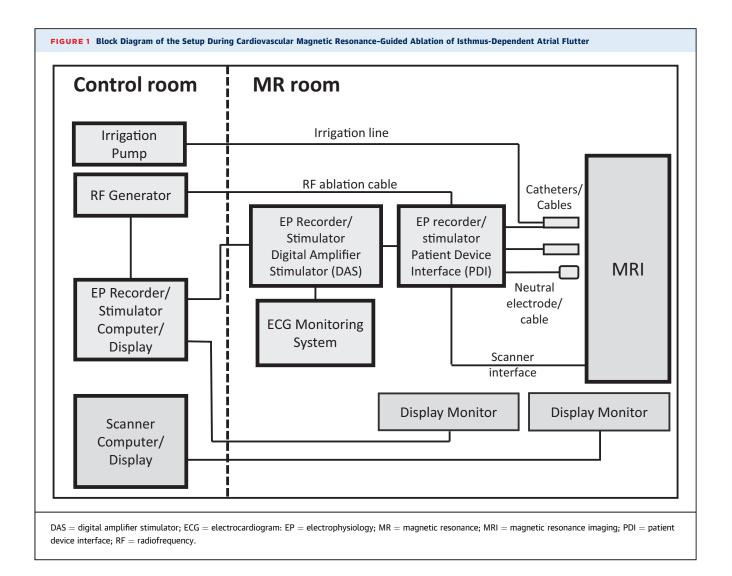
The aim of this study was to determine the safety and accuracy of this technique at 2-year follow-up.

METHODS

POPULATION. Between July 2021 and February 2022, consecutive patients with suspected CTI-dependent atrial flutter on electrocardiogram (ECG) were prospectively screened at our institution to undergo their first flutter ablation procedure using iCMR. Inclusion criteria were as follows: >18 years of age, suspected CTI-dependent atrial flutter on ECG, and signed informed consent. Exclusion criteria were a history of CTI-dependent flutter ablation, and contraindications to CMR (cerebral clip, metallic eye implant, and claustrophobia). All patients gave informed written consent for iCMR-guided flutter ablation and enrollment in the clinical research study. The study was approved by the local ethics committee of our institution and conducted in accordance with the Declaration of Helsinki. This study followed the Strengthening the Reporting of Observational Studies in Epidemiology reporting guideline for cohort studies.

ABLATION PROTOCOL. The principles of the electrophysiology protocol under iCMR have been previously described in details.¹⁻³ Briefly, the iCMR setup

Safety, Efficacy And 2-Year Outcomes Of Cardiovascular Magnetic Resonance-Guided Ablation Of Isthmus-Dependent Atrial Flutter Two Year Follow Up 1008 808 6008 408 208 0 Menth 12 Month 18 Month 24 Month Free from of Isthmus-Dependent Atrial Flutter



was similar to that previously described in previous studies.¹⁻⁴ The procedure was performed in a large bore 1.5-T MAGNETOM Aera Scanner (Siemens Healthineers) with 2 magnetic resonance-conditional, open-irrigated, steerable ablation catheters (VISION-MR; Imricor Medical Systems). The catheters were equipped with fiber optical temperature sensors and integrated MR tracking coils. They were connected to an electrophysiology workstation (Advantage-MR; Imricor Medical Systems) that provided monitoring and pacing capabilities, and to a radiofrequency (RF) generator operated from outside the CMR suite (Figure 1).

Patients were prepped on a dedicated and dockable CMR table in a preparation room just outside of the CMR suite. Patients were placed under conscious sedation. Midazolam (2 mg) was injected intravenously at the time of the venous puncture. Then,

sufentanil (5 μ g) was administered for analgesia. Two 9-F introducers were placed in the femoral vein under echography guidance, and patients were transferred to the CMR suite for flutter ablation.

Oxygen saturation and blood pressure were continuously monitored during the procedure in all patients. Catheters were tracked using receiver coils placed at the tip of the catheter under compressed sensing real-time cine CMR imaging (Video 1). The diagnostic catheter was introduced into the coronary sinus to serve as a reference catheter, and an irrigated ablation catheter was introduced in the right atrium to perform ablation shots at the CTI (the power of each shot was 50 W during 60 seconds). RF ablation was performed using a point-by-point method across the CTI until the block was confirmed.

PROCEDURAL SUCCESS. Procedural success was defined as the achievement of a bidirectional

TABLE 1 Patient Characteristics (N = 13)	
Age, y	76 (73-78)
Male	8 (62)
Heart disease	
None	6 (46)
Ischemic	2 (15)
Valvular heart disease	4 (31)
Myocarditis	1 (8)
LVEF, %	60 (60-60)
Antiarrhythmic drugs	
None	3 (23)
Beta-blockers	5 (38)
Amiodarone	5 (38)
Flecainide	3 (23)
Values are median (IQR) or n (%). LVEF = left ventricular ejection fraction.	

conduction block across the CTI. Then, patients were monitored in a dedicated room during at least 4 hours after the procedure. Patients were discharged on the same day after a follow-up ECG and in the absence of focal venous complications.

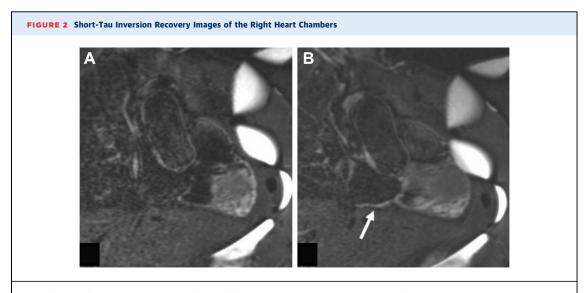
FOLLOW-UP. Patients were followed-up at 1 and 2 years by a clinical visit and an ECG. In case of palpitations, a 24-hour Holter ECG was performed.

STATISTICAL ANALYSIS. The data are presented as mean \pm SD and median (IQR) depending on the variable distribution. Descriptive statistical analyses were performed using PRISM 6 (GraphPad Software Inc).

RESULTS

Patients' characteristics are given in **Table 1**. Overall, 13 patients were included. The median age was 76 years (IQR: 73-78, range: 49-90). Among them, 1 patient had a history of myocarditis, 4 had valvular heart disease, 2 had ischemic heart disease, and 6 had no known heart disease (**Table 1**). All patients had a left ventricular ejection fraction >50%. All patients were under oral anticoagulant therapy for >1 month, and 10 patients received at least 1 antiarrhythmic drug. Eleven patients had atrial flutter during the procedure.

PROCEDURAL DATA. Ablation success was achieved in 11 patients (85%), as confirmed by the presence of a bidirectional conduction block across the CTI. The CTI ablation had to be completed in a conventional electrophysiology laboratory in 2 patients: 1 had severe chest pain during RF shots and needed general anesthesia, and the other deteriorated into atrial fibrillation during the procedure.



(A) Before radiofrequency ablation and (B) after radiofrequency ablation, showing an acute lesion of the cavotricuspid isthmus (hypersignal, arrow).

Bialobroda et al

No procedural or periprocedural complications (pericardial effusion or tamponade, vascular complications, or thromboembolic events) occurred.

In all patients, T2-weighted short-tau inversion recovery sequences centered on the CTI were acquired before and after ablation. All patients showed a clear hypersignal of the isthmus after ablation compared with baseline images, indicating the presence of focal myocardial edema (Figure 2). A prominent ridge was identified at the junction of the inferior vena cava and the right atrium in 1 patient (Figure 3).

FOLLOW-UP. Among 11 patients with procedural success, 9 (82%) had no recurrence of CTI-dependent atrial flutter at median follow-up of 27 months (IQR: 24-29) (Table 2). Two patients (18%) were lost to follow-up after ablation because they moved and were not reached despite several attempts.

DISCUSSION

This study demonstrates the accuracy and safety of iCMR-guided ablation of CTI-dependent atrial flutter at 2-year follow-up. Primary procedural success rate of iCMR-guided ablation was 85%, which is concordant with other studies.1 It is worth noting that 2 procedural failures were due either to sedation issues in the magnetic resonance suite, which can be overcome with proper anticipation, or to the need for electrical cardioversion. In the current study, no patient experienced recurrence of common atrial flutter during a 2-year follow-up. Nevertheless, the procedural success rate is lower than that with catheter ablation in an electrophysiology laboratory with fluoroscopy. The procedure was stopped in 1 patient because the flutter deteriorated into atrial fibrillation and required electrical cardioversion. A CMRcompatible defibrillation device could overcome this issue and would facilitate the procedures. In the magnetic resonance environment, a patient's pain is more difficult to manage than in a conventional electrophysiology laboratory, because of suite accessibility, the need of CMR compatible syringe pump, and appropriate ventilation support devices. Overall, no complication occurred, and all procedures were performed on an outpatient basis. Besides the absence of ionizing radiation, several advantages may be highlighted such as the visualization of the CTI before ablation. Indeed, the presence of a CTI ridge, the visualization of the Eustachian valve, and its

FIGURE 3 Identification of a Prominent Cavotricuspid Isthmus Ridge During Planning Cardiovascular Magnetic Resonance



The arrow indicates the cavotricuspid isthmus.

interaction with the CTI may help accelerate catheter access to the isthmus.

This technique also allows the visualization of post-RF endocardial damage through CMR tissue characterization. Bijvoet et al³ recently demonstrated that tissue characterization of acute lesions during CMR-guided CTI-dependent atrial flutter ablation demonstrates edema, perfusion defects, and necrosis with a core of microvascular damage. It can be assumed that these lesions progress to fibrosis and focal scar; however, in this study, CMR follow-up was not performed at 2 years.

TABLE 2 Cavotricuspid Isthmus Ablation Parameters and Follow-Up	
Interventional procedure duration, min	60 (60-120)
Total radiofrequency pulse duration, s	478 (206-577)
Obtention of bidirectional isthmus block, %	11 (85)
Double potentials, ms	90 (77-120)
Procedural complications	
Pericardial effusion/tamponade	0 (0)
Vascular complications	0 (0)
Thromboembolic events	0 (0)
Outpatients	13 (100)
Lost to follow-up	2 (18)
Follow-up, mo	27 (24-29)
Recurrence of isthmus-dependent atrial flutter	0 (0)

safety and efficacy of RF ablation of VT associated with ischemic cardiomyopathy using the Vision-MR

The main limitations of the study are as follows. First, although endocardium, myocardium, cardiac chambers, and structures are remarkably depicted in real time, spatial resolution is lower and real-time tissue characterization is not available. Second, active catheter tracking is accurate but at a lower temporal resolution than in fluoroscopy. Third, significant interferences occur between the CMR field and the electrograms, requiring stepwise point-bypoint ablation. Fourth, significant interferences may occur between RF and electrograms. Finally, the CMR environment requires technical adaptations (wires between the CMR suite and the pump, RF generator, EP recording system). Well-established safety and emergency procedures are needed and derived from stress CMR studies.

iCMR flutter ablation may pave the way to future ventricular tachycardia (VT) ablation procedures in the magnetic resonance environment. iCMR-guided VT ablation would enable RF ablation on scarred tissue without the need to induce VT and with direct visualization of myocardial substrates. The feasibility of CMR guidance and real-time assessment of ablation injury has been demonstrated in the porcine left ventricle. The VISABL-VT (Vision-MR Ablation Catheter 2.0 for the Treatment of Ventricular Tachycardia) trial is a prospective, single-arm, multicenter, interventional study investigating the

CONCLUSIONS

(NCT05543798).

iCMR-guided ablation of CTI-dependent atrial flutter demonstrates a good safety profile and efficacy at 2-year follow-up.

Ablation Catheter 2.0 in the iCMR environment

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The authors have reported that they have no relationships relevant to the contents of this paper to disclose.

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KEY WORDS CMR-guided ablation, interventional cardiovascular magnetic resonance, outcomes, procedural success, radiofrequency atrial flutter ablation

APPENDIX For supplemental videos, please see the online version of this paper.